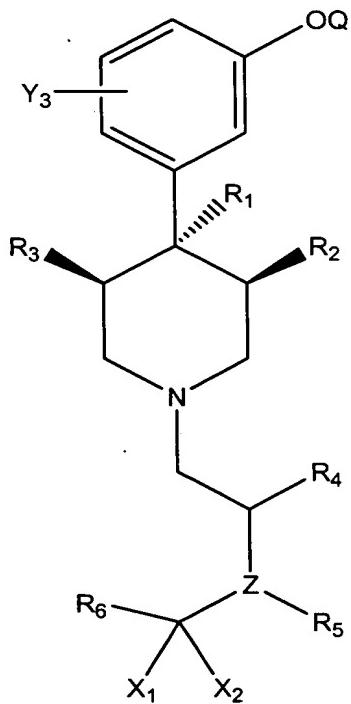


CLAIMS

Please amend the claims as follows:

Claim 1 (Currently Amended): A method of binding a kappa opioid receptor in a subject in need thereof, comprising:

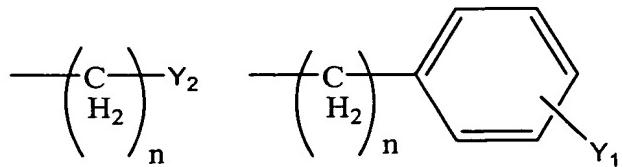
administering to said subject a composition comprising a kappa opioid receptor antagonist and a physiologically acceptable carrier, wherein the kappa opioid receptor antagonist is a compound of formula (1):

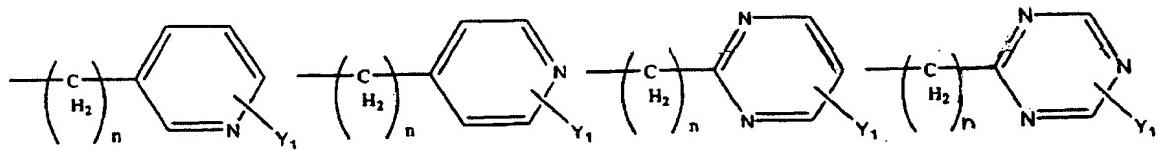


(I)

wherein Q is H or COC₁₋₈ alkyl;

R₁ is C₁₋₈ alkyl, or one of the following structures:





Y_1 , is H, OH, Br, Cl, F, CN, CF_3 , NO_2 , N_3 , OR_8 , CO_2R_9 , C_{1-6} alkyl, $NR_{10}R_{11}$, $NHCOR_{12}$, NHO_2R_{12} , $CONR_{13}R_{14}$, or $CH_2(CH_2)_nY_2$;

Y_2 is H, CF_3 , CO_2R_9 , C_{1-6} alkyl, $NR_{10}R_{11}$, $NHCOR_{12}$, NHO_2R_{12} , $CONR_{13}R_{14}$, CH_2OH , CH_2OR_8 , or $COCH_2R_9$;

Y_3 is H, OH, Br, Cl, F, CN, CF_3 , NO_2 , N_3 , OR_8 , CO_2R_9 , C_{1-6} alkyl, $NR_{10}R_{11}$, $NHCOR_{12}$, NHO_2R_{12} , $CONR_{13}R_{14}$, or $CH_2(CH_2)_nY_2$;

R_2 is H, C_{1-8} alkyl, C_{3-8} alkenyl, C_{3-8} alkynyl or CH_2 aryl substituted by one or more groups Y_1 ;

R_3 is H, C_{1-8} alkyl, C_{3-8} alkenyl, C_{3-8} alkynyl or CH_2 aryl substituted by one or more groups Y_1 ;

wherein R_2 and R_3 may be bonded together to form a C_{2-8} , alkyl group;

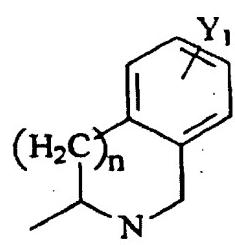
R_4 is hydrogen, C_{1-8} alkyl, CO_2C_{1-8} alkylaryl substituted by one or more groups Y_1 , CH_2 aryl substituted by one or more groups Y_1 or CO_2C_{1-8} alkyl;

Z is N, O or S; where Z is O or S, there is no R_5

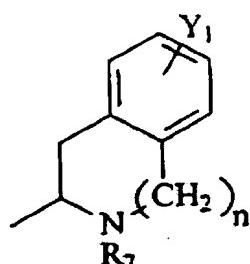
R_5 is H, C_{1-8} alkyl, C_{3-8} alkenyl, C_{3-8} alkynyl, $CH_2CO_2C_{1-8}$ alkyl, CO_2C_{1-8} alkyl or CH_2 aryl substituted by one or more groups Y_1 ;

n is 0, 1, 2 or 3;

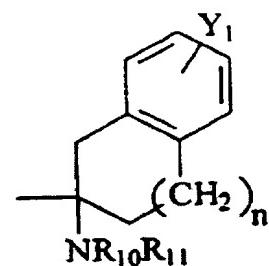
R_6 is a group selected from the group consisting of structures (a)-(w) and (cc)-(bbb):



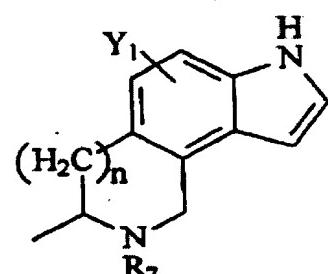
(a)



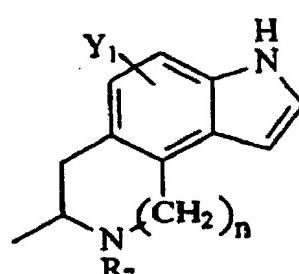
(b)



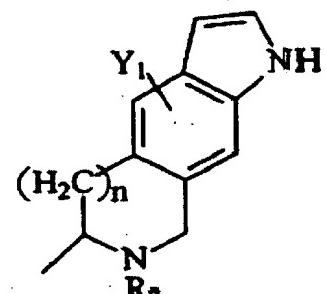
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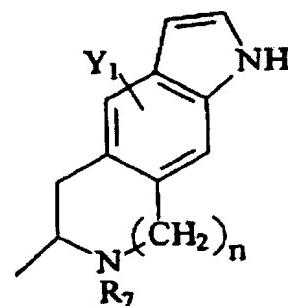
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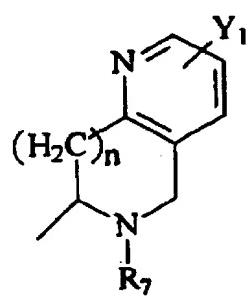
(e)



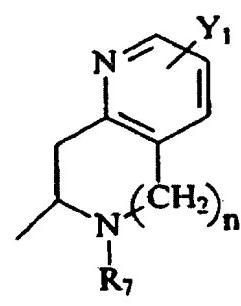
(f)



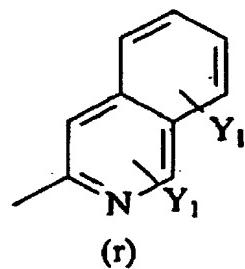
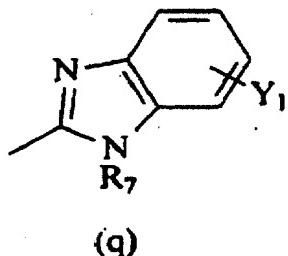
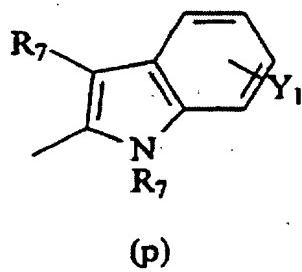
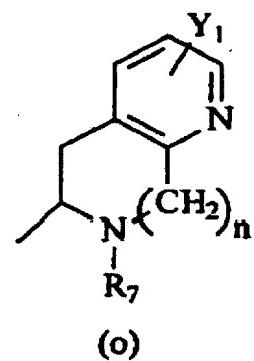
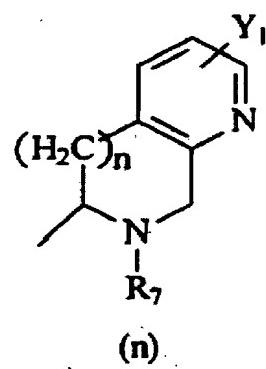
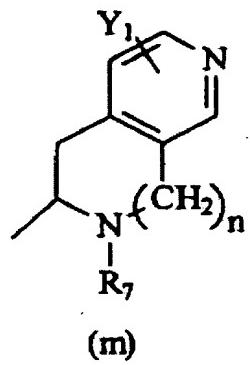
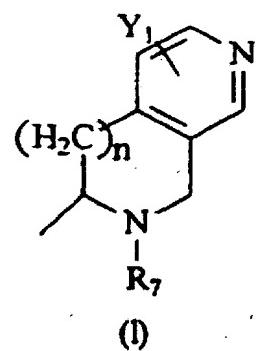
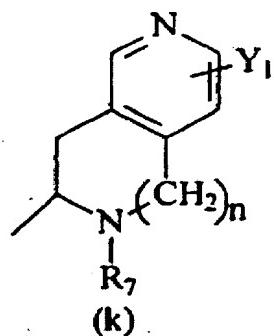
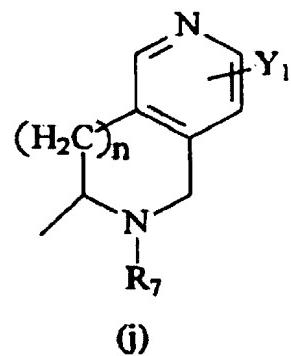
(g)

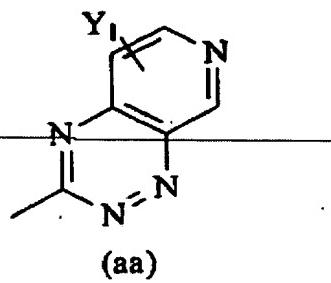
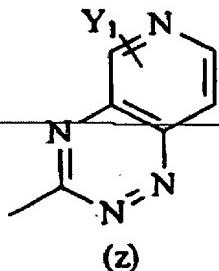
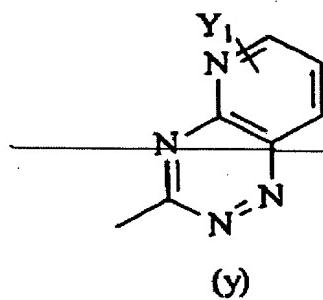
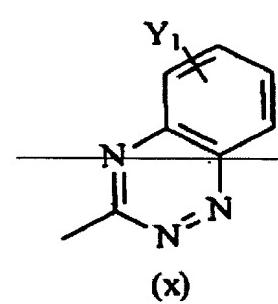
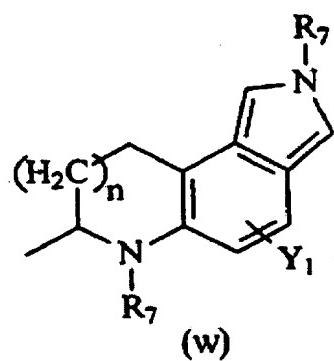
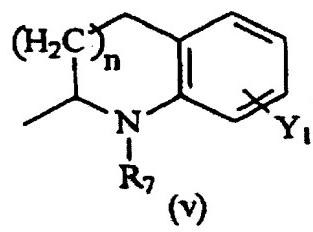
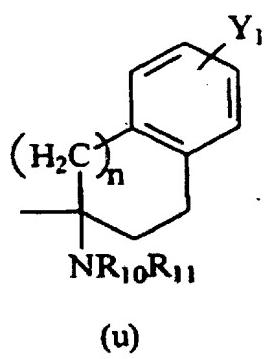
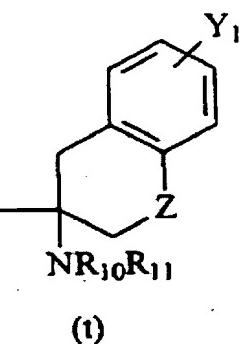
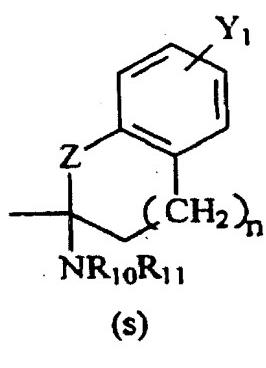


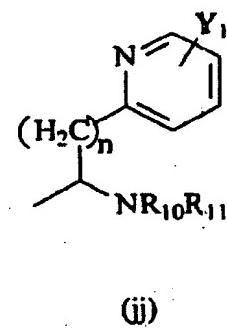
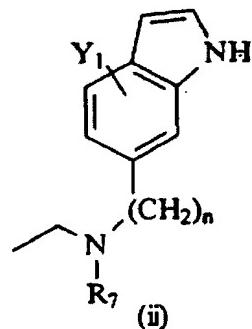
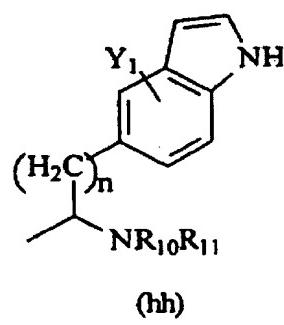
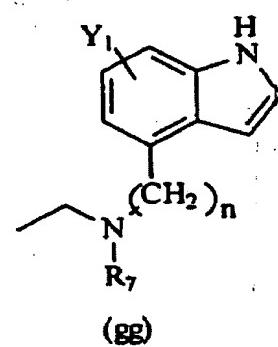
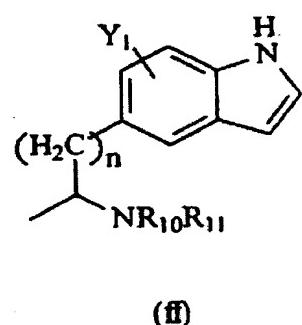
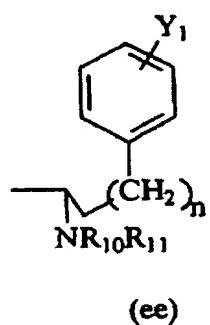
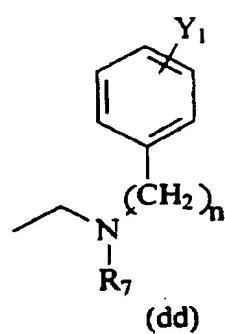
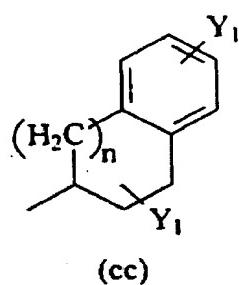
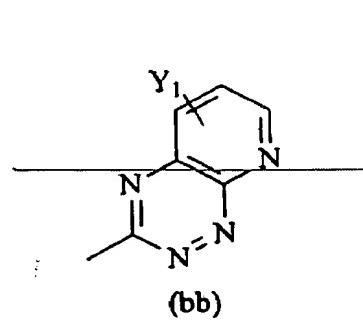
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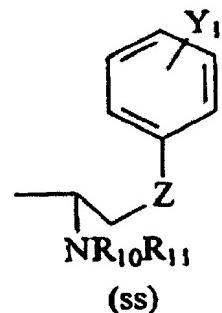
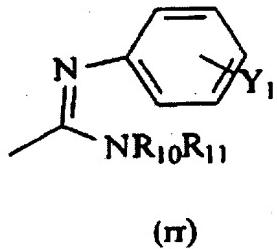
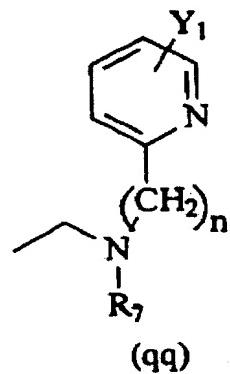
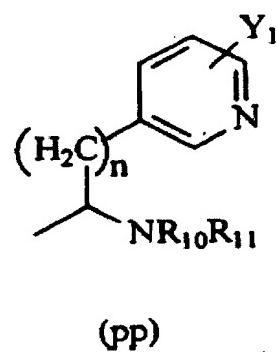
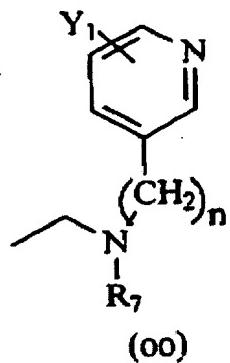
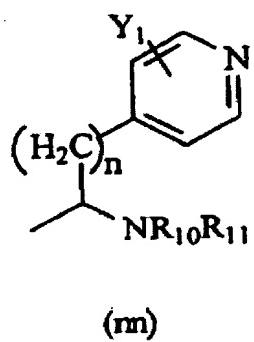
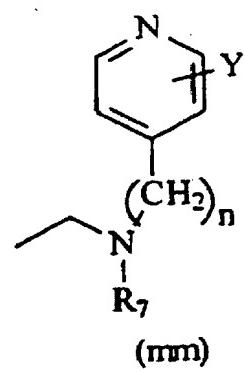
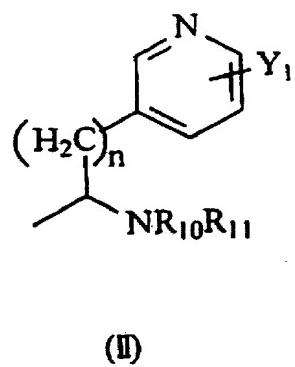
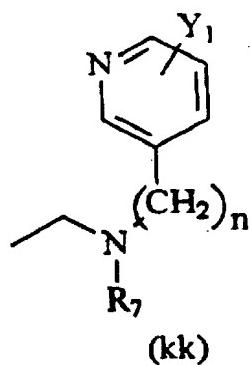


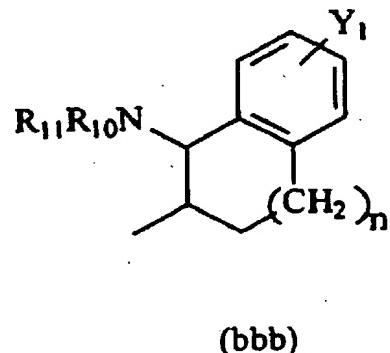
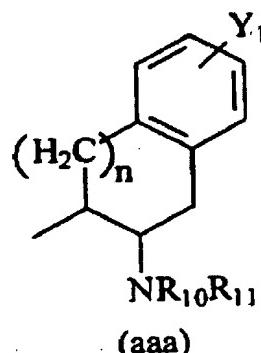
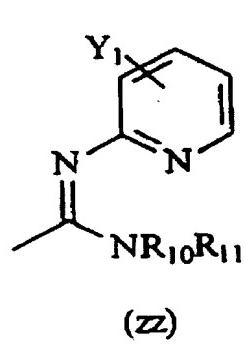
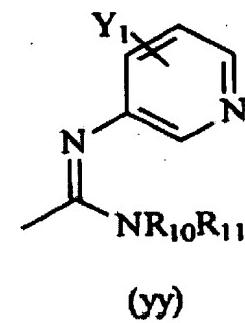
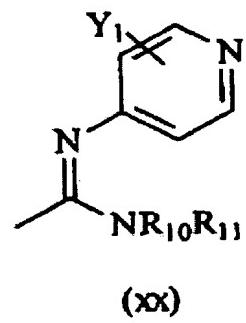
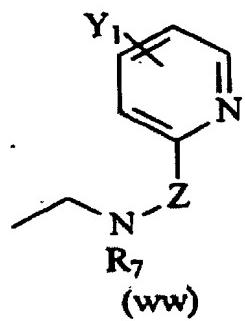
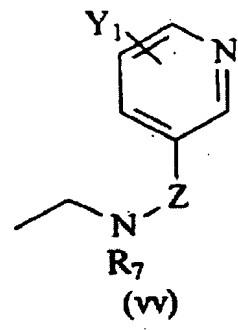
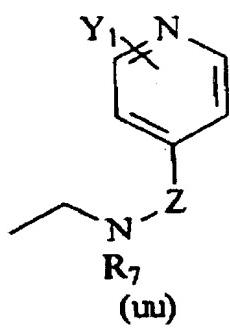
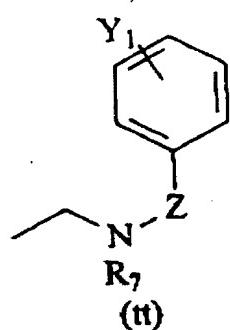
(i)











X₁ is hydrogen, C₁₋₈ alkyl, C₃₋₈alkenyl, or C₃₋₈alkynyl;

X₂ is hydrogen, C₁₋₈alkyl, C₃₋₈alkenyl, or C₃₋₈alkynyl; or

X₁ and X₂ together form =O, =S, =NH;

R₇ is H, C₁₋₈alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀, R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, CH₂(CH₂)_nY₂, or C(=NH)NR₁₆R₁₇;

R₈ is H, C₁₋₈alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl;

R₉ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl;

R₁₀ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl ;

R₁₁ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl ;

R₁₂ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl ;

R₁₃ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl ;

R₁₄ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl ;

R₁₅ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl ;

R₁₆ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl ; and

R₁₇ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆alkyl;

and pharmaceutically acceptable salts thereof.

Claim 2 (Currently Amended): The method of claim 1, wherein said kappa opioid receptor antagonist is a compound of formula (I), wherein R₁, R₄, R₅, Y₁, Y₂, Z, n, X₁, X₂, and R₇-R₁₇ are as in Claim 1;

Y₃ is H;

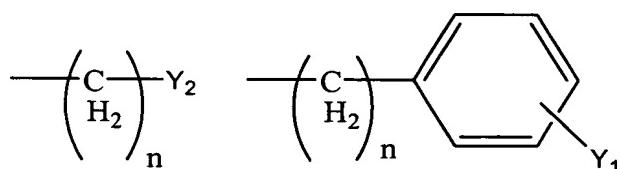
R₂ and R₃ are each, independently, H, C₁₋₈ alkyl, C₃₋₈ alkynyl, C₃₋₈ alkynyl, or CH₂aryl substituted by one or more substituents Y₁; and

R₆ is a group having a formula selected from the group consisting of structures (a)-(w) and (cc);

and pharmaceutically acceptable salts thereof.

3. (Previously Presented) The method of claim 1, wherein said kappa opioid receptor antagonist is a compound of formula (I) wherein Y₁, Y₂, R₄, R₅, Z, n, X₁, X₂ and R₈-R₁₅ are as in Claim 1;

R₁ is C₁₋₈ alkyl, or one of the following structures:



Y₃ is H;

R₂ and R₃ are each, independently, H or C₁₋₈ alkyl, wherein R₂ and R₃ cannot both be H at the same time;

R₆ is a formula selected from the structures (a)-(r); and

R₇ is H, C₁₋₈ alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, or CH₂(CH₂)_nY₂.

Claim 4 (Previously Presented) The method of claim 1, wherein said kappa opioid receptor antagonist is a compound of formula (I) wherein Y₁, Z, n, X₁, X₂ and R₈-R₁₅ are as in Claim 1;

R₁ is C₁₋₈ alkyl;

Y₂ is H, CF₃, CO₂R₉, C₁₋₆ alkyl, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, CH₂OH, CH₂OR₈, or COCH₂R₉;

Y₃ is H;

R₂ and R₃ are each, independently, H or methyl, wherein R₂ and R₃ cannot both be H at the same time;

R₄ is H, C₁₋₈ alkyl, CO₂C₁₋₈alkyl, or CH₂ aryl substituted by one or more substituents Y₁ and the stereocenter adjacent to R₄ is in an (S) configuration;

R₅ is H, C₁₋₈ alkyl, or CH₂CO₂C₁₋₈ alkyl;

R₆ is a group having a formula selected from the group consisting of structures (a)-(c) and (h)-(o); and

R₇ is H, C₁₋₈alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, or CH₂(CH₂)_nY₂.

Claim 5 (Previously Presented) The method of claim 1, wherein said kappa opioid receptor antagonist is a compound of formula (I), wherein Y₁, Z, n, X₁, X₂ and R₈-R₁₄ are as in Claim 1;

R₁ is methyl,

Y₂ is H, CF₃, CO₂R₉, C₁₋₆ alkyl, NR₁₀R₁₁, NHCO₂R₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, CH₂OH, CH₂OR₈, or COCH₂R₉;

Y₃ is H;

R₂ and R₃ are each H or methyl, such that when R₂ is H, R₃ is methyl and vice versa;

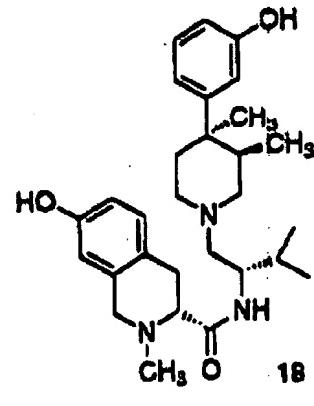
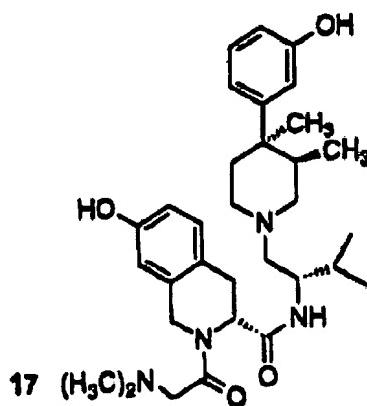
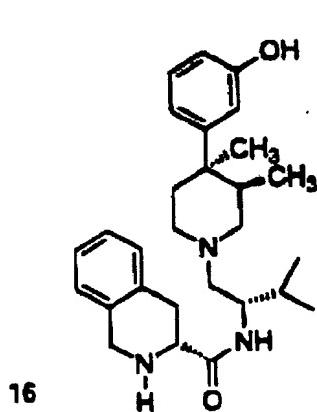
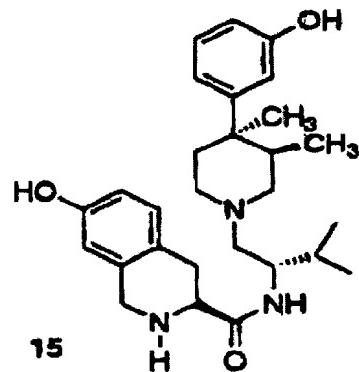
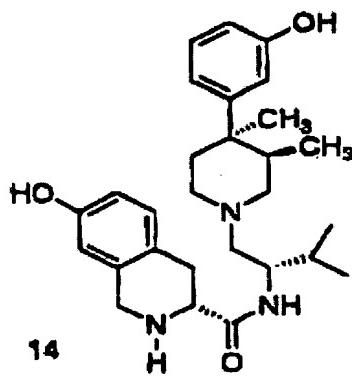
R₄ is C₁₋₈ alkyl, or CO₂C₁₋₈ alkyl, and the stereocenter adjacent to R₄ has a configuration of (S);

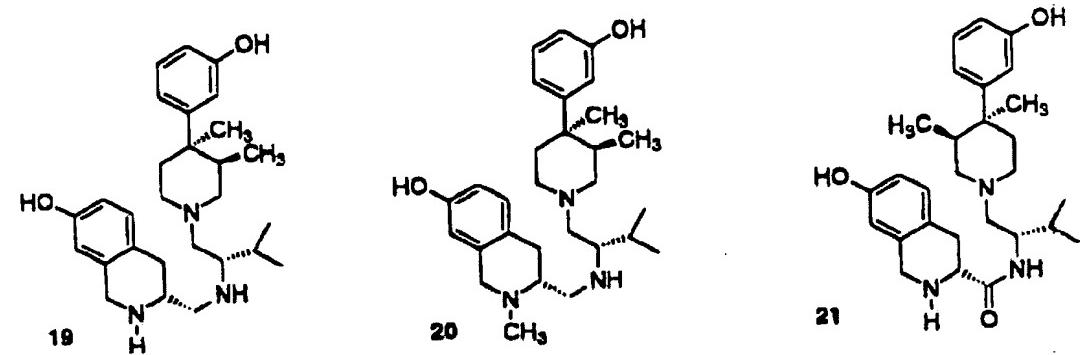
R₅ is H;

R₆ is a group having a formula selected from the group consisting of structures (a) and (b); and

R₇ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents Y₁, or CH₂(CH₂)_nY₂.

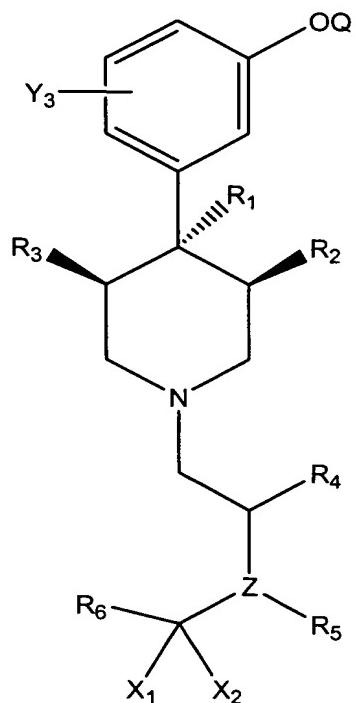
Claim 6 (Currently Amended) The method of claim 1, wherein said kappa opioid receptor antagonist is a compound selected from formulae 14-21 ~~of Fig. 1~~ as follows:





Claim 7 (Currently Amended) A kappa opioid receptor antagonist compound

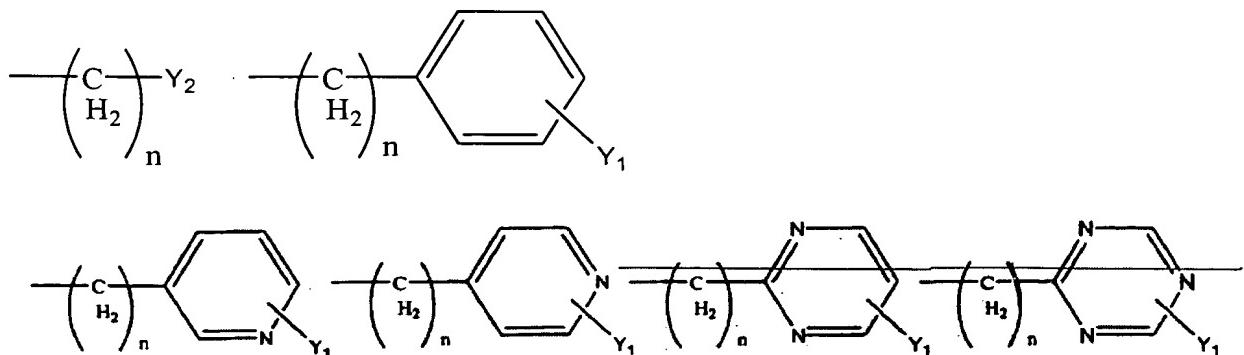
represented by the formula (I):



(I)

wherein Q is H or COC_{1-8} alkyl;

R_1 is C_{1-8} alkyl, or one of the following structures:



Y_1 is H, OH, Br, Cl, F, CN, CF_3 , NO_2 , N_3 , OR_8 , CO_2R_9 , C_{1-6} alkyl, $\text{NR}_{10}\text{R}_{11}$,

NHCOR_{12} , $\text{NHCO}_2\text{R}_{12}$, $\text{CONR}_{13}\text{R}_{14}$, or $\text{CH}_2(\text{CH}_2)_n\text{Y}_2$;

Y_2 is H, CF_3 , CO_2R_9 , C_{1-6} alkyl, $\text{NR}_{10}\text{R}_{11}$, NHCOR_{12} , $\text{NHCO}_2\text{R}_{12}$, $\text{CONR}_3\text{R}_{14}$,

CH_2OH , CH_2OR_8 , or COCH_2R_9 ;

Y_3 is H, OH, Br, Cl, F, CN, CF_3 , NO_2 , N_3 , OR_8 , CO_2R_9 , C_{1-6} alkyl, $\text{NR}_{10}\text{R}_{11}$,

NHCOR_{12} , $\text{NHCO}_2\text{R}_{12}$, $\text{CONR}_{13}\text{R}_{14}$, or $\text{CH}_2(\text{CH}_2)_n\text{Y}_2$;

R_2 is H, C_{1-8} alkyl, C_{3-8} alkenyl, C_{3-8} alkynyl or CH_2aryl substituted by one or more groups Y_1 ;

R_3 is H, C_{1-8} alkyl, C_{3-8} alkenyl, C_{3-8} alkynyl or CH_2aryl substituted by one or more groups Y_1 ;

wherein R_2 and R_3 may be bonded together to form a C_{2-8} alkyl group;

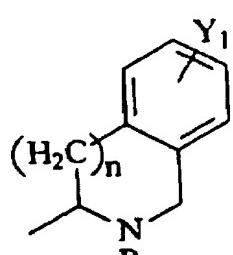
R_4 is hydrogen, C_{1-8} alkyl, $\text{CO}_2\text{C}_{1-8}$ alkylaryl substituted by one or more groups Y_1 , CH_2aryl substituted by one or more groups Y_1 or $\text{CO}_2\text{C}_{1-8}$ alkyl;

Z is N, O or S; when Z is O or S there is no R_5

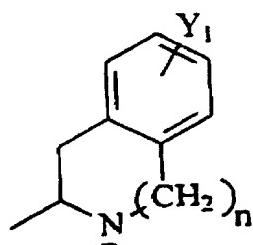
R_5 is H, C_{1-8} alkyl, C_{3-8} alkenyl, C_{3-8} alkynyl, $\text{CH}_2\text{CO}_2\text{C}_{1-8}$ alkyl, $\text{CO}_2\text{C}_{1-8}$ alkyl or CH_2aryl substituted by one or more groups Y_1 ;

n is 0, 1, 2 or 3;

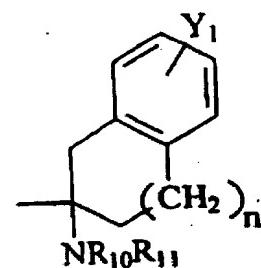
R_6 is a group selected from the group consisting of structures (a)-(w) and (cc)-(bbb):



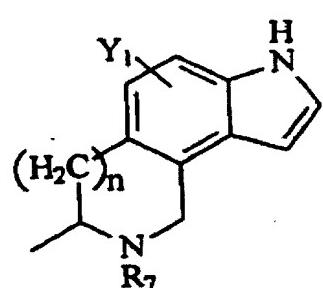
(a)



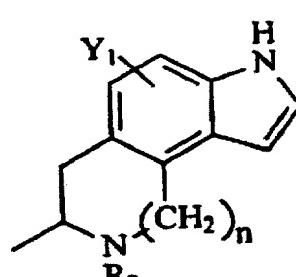
(b)



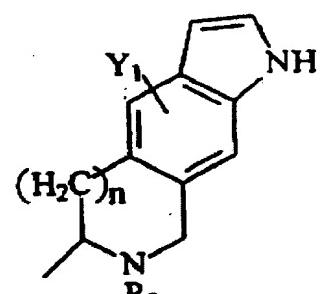
(c)



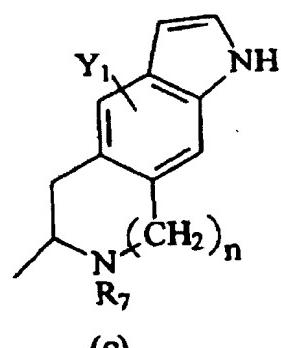
(d)



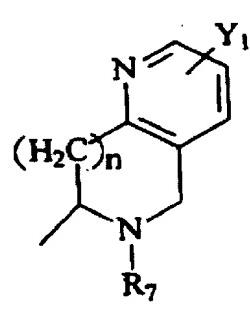
(e)



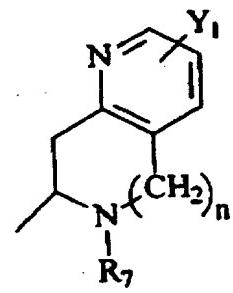
(f)



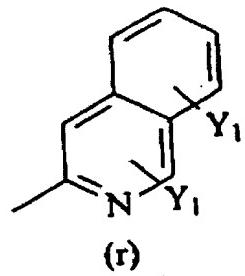
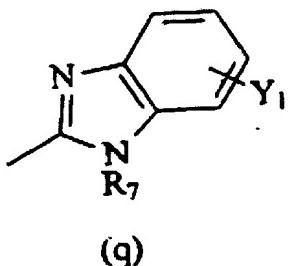
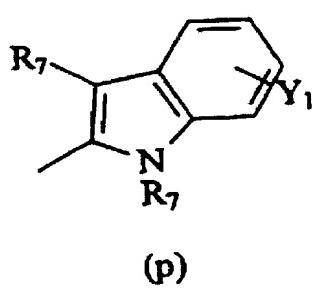
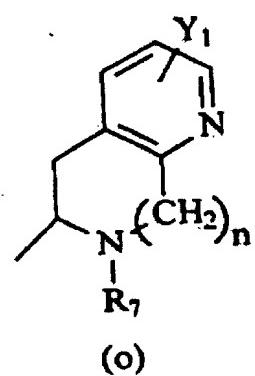
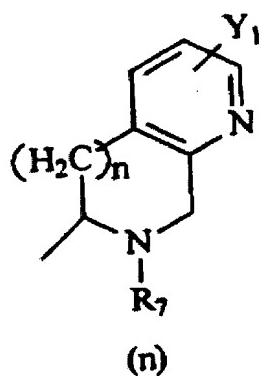
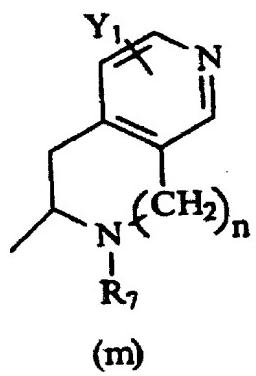
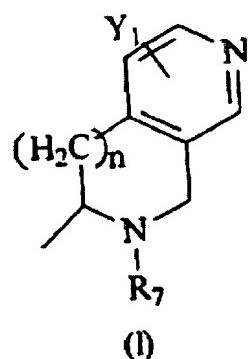
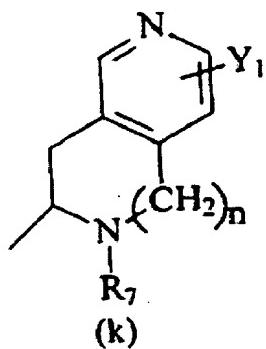
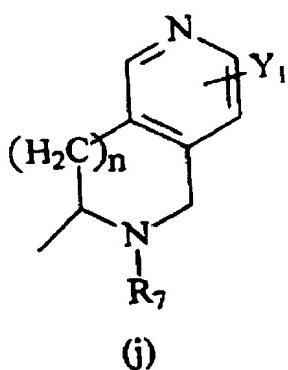
(g)

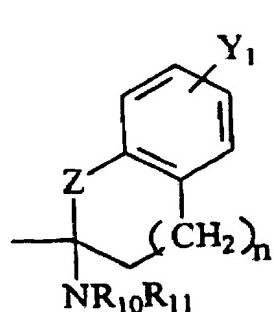


(h)

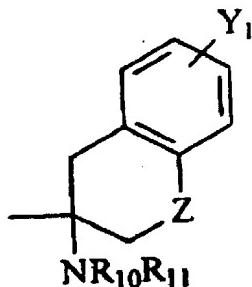


(i)

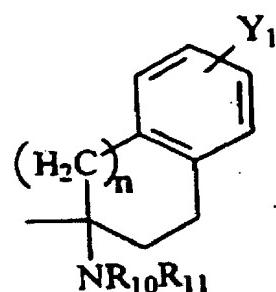




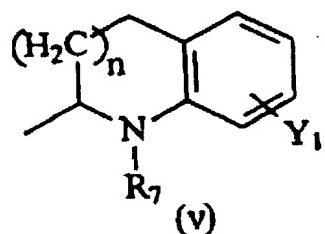
(s)



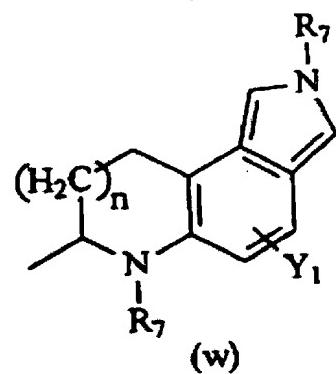
(t)



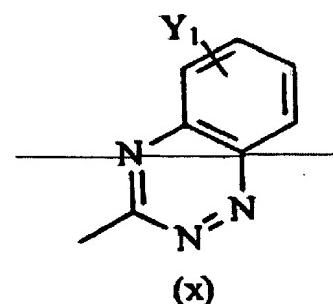
(u)



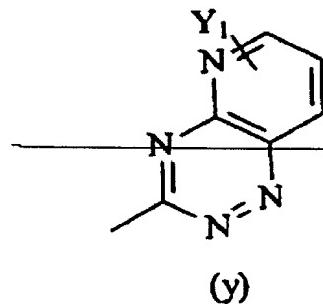
(v)



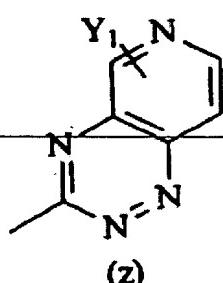
(w)



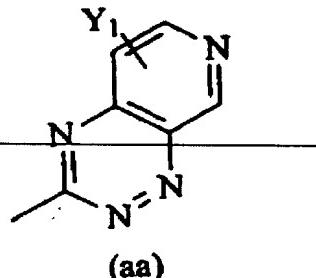
(x)



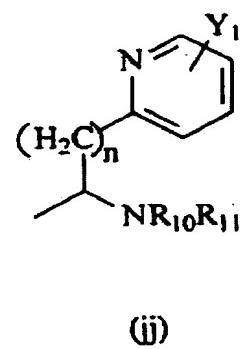
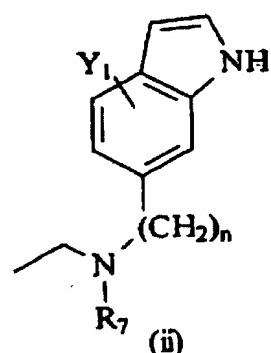
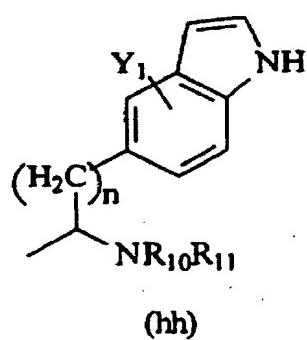
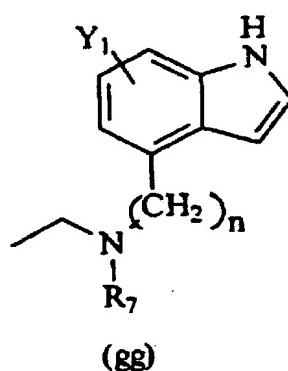
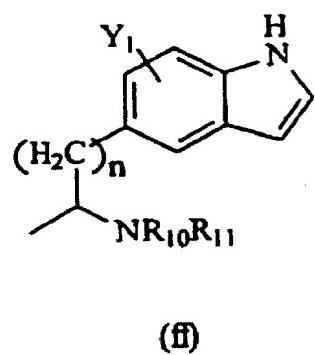
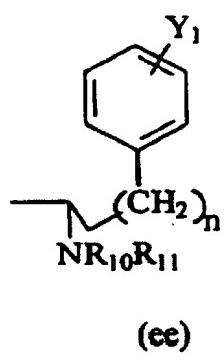
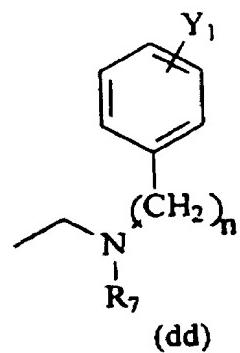
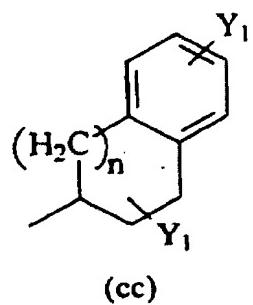
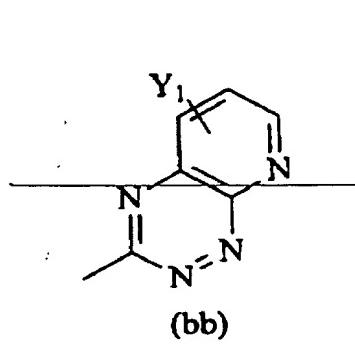
(y)

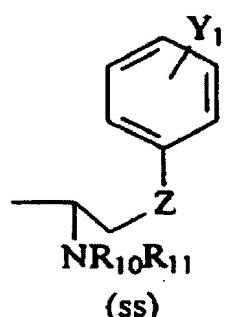
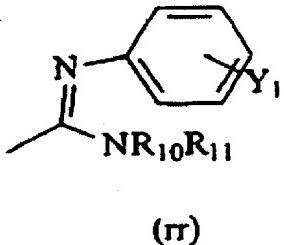
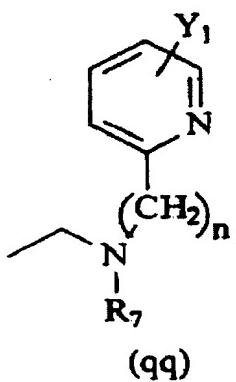
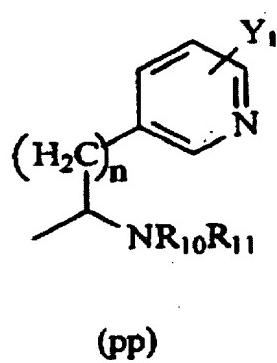
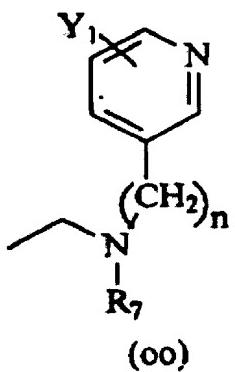
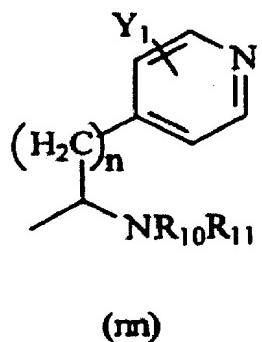
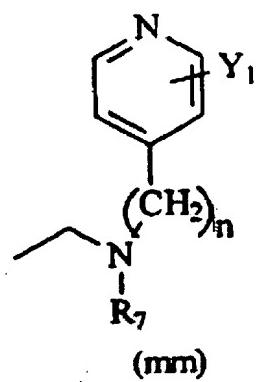
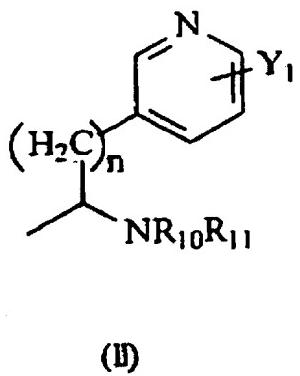
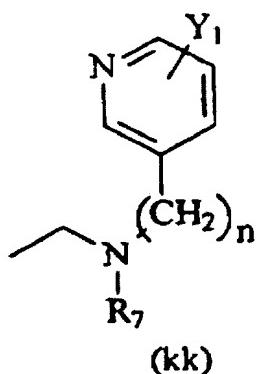


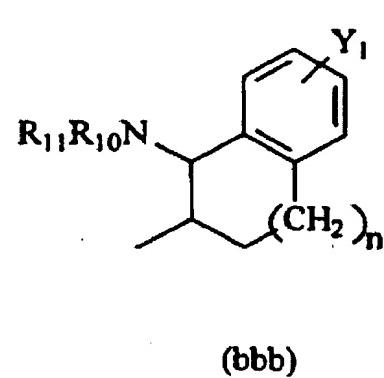
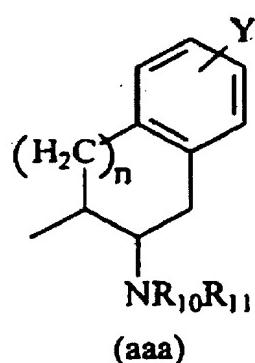
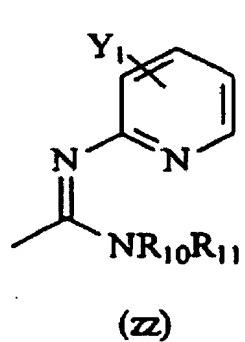
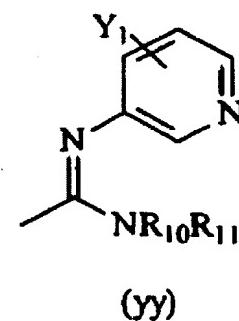
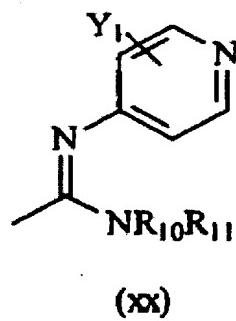
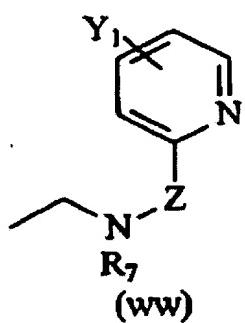
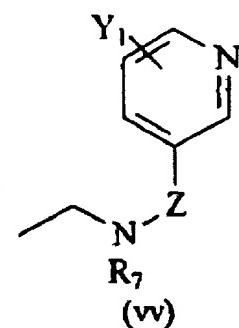
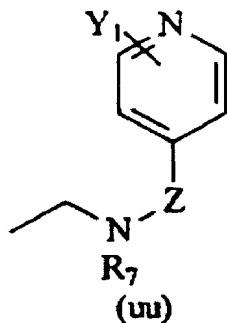
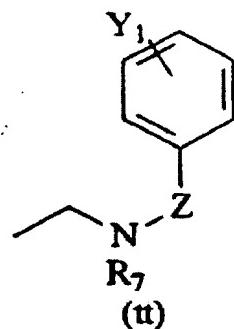
(z)



(aa)







X₁ is hydrogen, C₁₋₈ alkyl, C₃₋₈ alkenyl, or C₃₋₈ alkynyl;

X₂ is hydrogen, C₁₋₈ alkyl, C₃₋₈ alkenyl, or C₃₋₈ alkynyl;

or X₁ and X₂ together form =O, =S, or =NH;

R₇ is H, C₁₋₈ alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, CH₂(CH₂)_nY₂, or C(=NH)NR₁₆R₁₇;

R₈ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl;

R₉ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl;

R₁₀ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl ;

R₁₁ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl ;

R₁₂ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl ;

R₁₃ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl ;

R₁₄ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl ;

R₁₅ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl ;

R₁₆ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl ; and

R₁₇ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY_{2'}; wherein Y_{2'} is H, CF₃, or C₁₋₆ alkyl

and pharmaceutically acceptable salts thereof.

Claim 8 (Currently Amended) The kappa opioid receptor antagonist compound of claim 7, wherein R₁, R₄, R₅, Y₁, Y₂, Z, n, X₁, X₂, and R₇-R₁₇ are as in Claim 7;

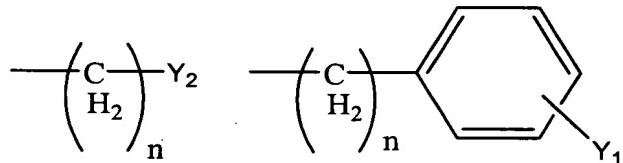
Y₃ is H;

R₂ and R₃ are each, independently, H, C₁₋₈ alkyl, C₃₋₈ alkynyl, C₃₋₈ alkynyl, or CH₂aryl substituted by one or more substituents Y₁; and

R₆ is a group having a formula selected from the group consisting of structures (a)-(w) and (cc).

Claim 9 (Previously Presented) The kappa opioid receptor antagonist compound of claim 7, wherein Y₁, Y₂, R₄, R₅, Z, n, X₁, X₂ and R₈R₁₅ are as in Claim 7;

R₁ is C₁₋₈ alkyl, or one of the following structures:



Y₃ is H;

R₂ and R₃ are each, independently, H or C₁₋₈ alkyl, wherein R₂ and R₃ cannot both be H at the same time;

R₆ is a formula selected from the structures (a)-(r); and

R₇ is H, C₁₋₈ alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, or CH₂(CH₂)_nY₂.

Claim 10 (Previously Presented) The kappa opioid receptor antagonist compound of claim 7, wherein Y₁, Z, n, X₁, X₂ and R₈-R₁₅ are as in Claim 7;

R₁ is C₁₋₈ alkyl;

Y₂ is H, CF₃, CO₂R₉, C₁₋₆ alkyl, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, CH₂OH, CH₂OR₈, or COCH₂R₉;

Y₃ is H;

R₂ and R₃ are each, independently, H or methyl, wherein R₂ and R₃ cannot both be H at the same time;

R₄ is H, C₁₋₈ alkyl, CO₂C₁₋₈alkyl, or CH₂ aryl substituted by one or more substituents Y₁ and the stereocenter adjacent to R₄ is in an (S) configuration;

R₅ is H, C₁₋₈ alkyl, CH₂CO₂C₁₋₈ alkyl;

R₆ is a group having a formula selected from the group consisting of structures (a)-(c) and (h)-(o); and

R₇ is H, C₁₋₈alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, or CH₂(CH₂)_nY₂.

Claim 11 (Previously Presented) The kappa opioid receptor antagonist compound of claim 7, wherein Y₁, Z, n, X₁, X₂ and R₈-R₁₄ are as in Claim 7;

R₁ is methyl,

Y₂ is H, CF₃, CO₂R₉, C₁₋₆ alkyl, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, CH₂OH, CH₂OR₈, or COCH₂R₉;

Y₃ is H;

R₂ and R₃ are each H or methyl, such that when R₂ is H, R₃ is methyl and vice versa;

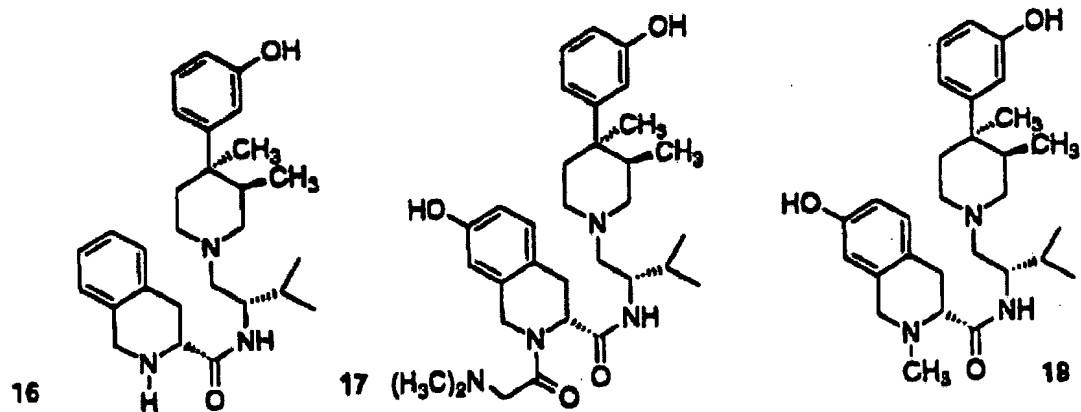
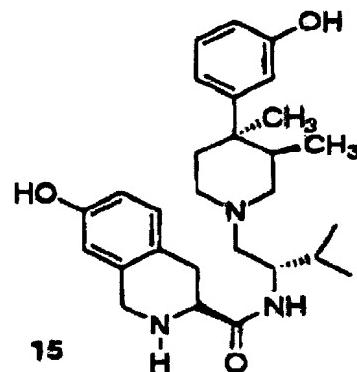
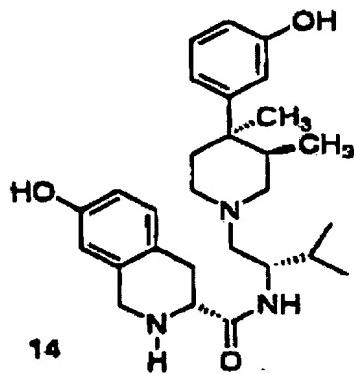
R₄ is C₁₋₈ alkyl, or CO₂C₂₋₈ alkyl, and the stereocenter adjacent to R₄ has a configuration of (S);

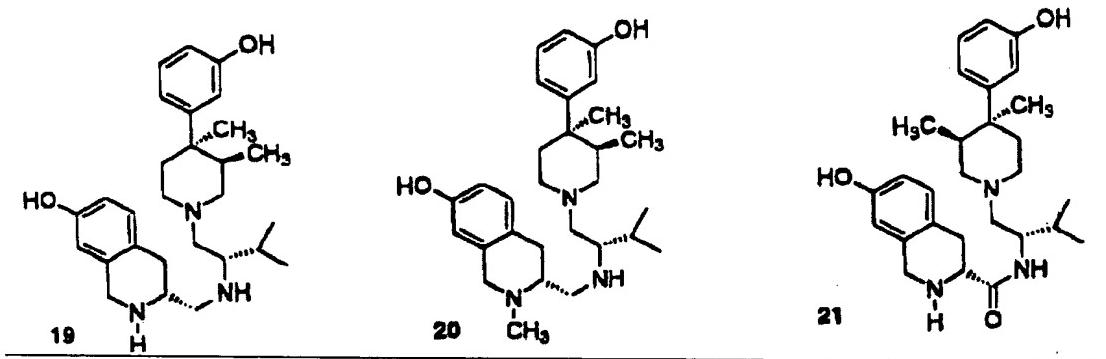
R₅ is H;

R₆ is a group having a formula selected from the group consisting of structures (a) and (b); and

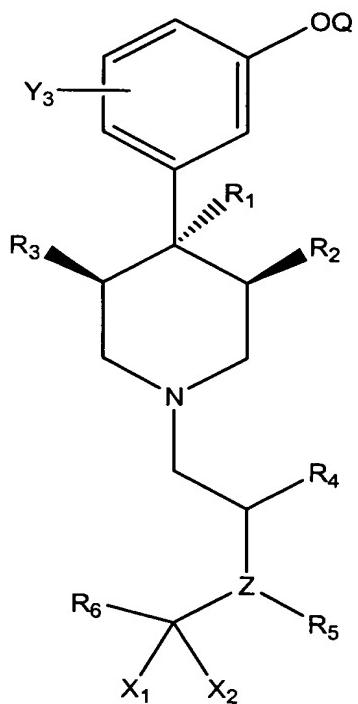
R₇ is H, C₁₋₈ alkyl, CH₂aryl substituted by one or more substituents Y₁ or CH₂(CH₂)_nY₂.

Claim 12 (Currently Amended) The kappa opioid receptor antagonist of claim 7, wherein said compound is a compound selected from formulae 14-21 of Fig. 1 as follows:





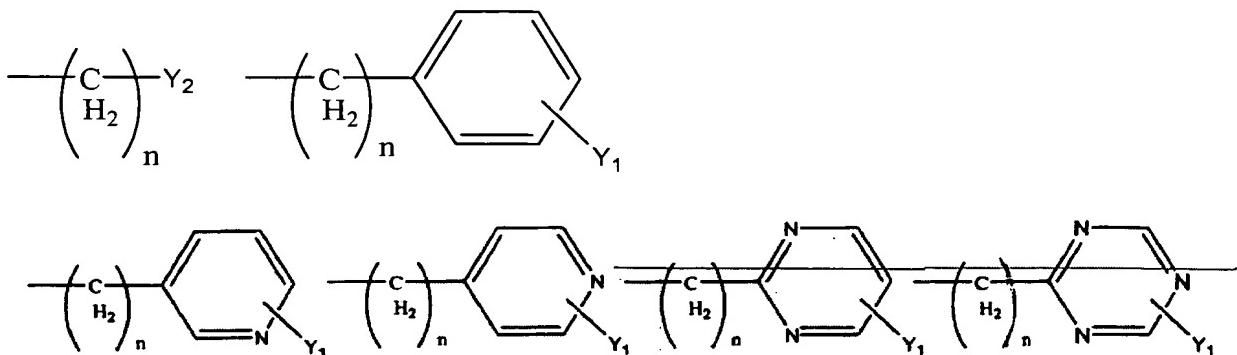
Claim 13 (Currently Amended) A pharmaceutical composition comprising:
an effective amount of a kappa opioid receptor antagonist and a physiologically acceptable carrier, wherein the kappa opioid receptor antagonist is a compound of formula (I):



(I)

wherein Q is H or COC₁₋₈ alkyl;

R₁ is C₁₋₈ alkyl, or one of the following structures:



Y₁ is H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, OR₈, CO₂R₆, C₁₋₆ alkyl, NR₁₀R₁₁,

NHCOR₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, or CH₂(CH₂)_nY₂;

Y₂ is H, CF₃, CO₂R₉, C₁₋₆alkyl, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₂, CONR₃R₁₄,

CH₂OH, CH₂OR₈, or COCH₂R₉;

Y₃ is H, OH, Br, Cl, F, CN, CF₃, NO₂, N₃, OR₈, CO₂R₉, C₁₋₆ alkyl, NR₁₀R₁₁,

NHCOR₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, or CH₂(CH₂)_nY₂;

R₂ is H, C₁₋₈ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl or CH₂aryl substituted by one or more groups Y₁;

R₃ is H, C₁₋₈ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl or CH₂aryl substituted by one or more groups Y₁;

wherein R₂ and R₃ may be bonded together to form a C₂₋₈ alkyl group;

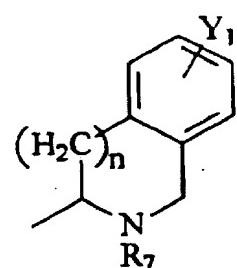
R₄ is hydrogen, C₁₋₈ alkyl, CO₂C₁₋₈ alkylaryl substituted by one or more groups Y₁, CH₂aryl substituted by one or more groups Y₁, or CO₂C₁₋₈ alkyl;

Z is N, O or S; when Z is O or S, there is no R₅

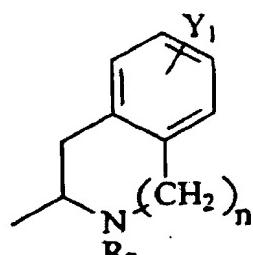
R₅ is H, C₁₋₈ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, CH₂CO₂C₁₋₈ alkyl, CO₂C₁₋₈ alkyl or CH₂aryl substituted by one or more groups Y₁;

n is 0, 1, 2 or 3;

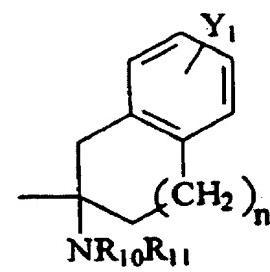
R₆ is a group selected from the group consisting of structures (a)-(w) and (cc)-(bbb):



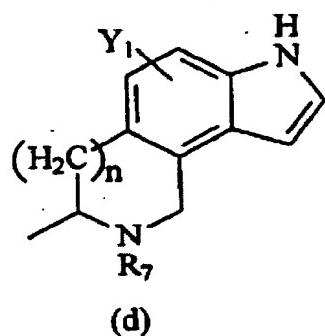
(a)



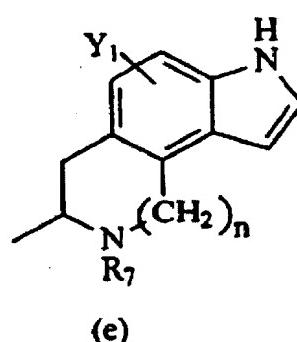
(b)



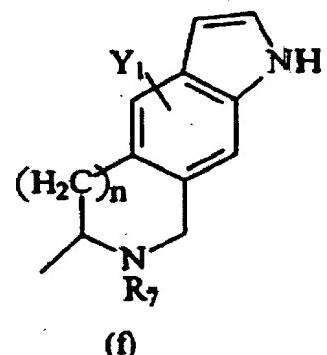
(c)



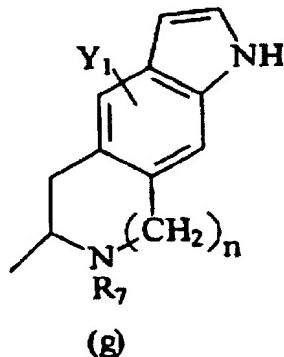
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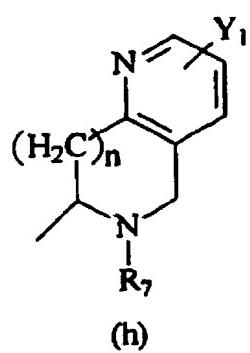
(e)



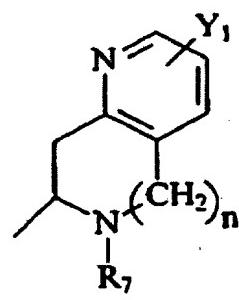
(f)



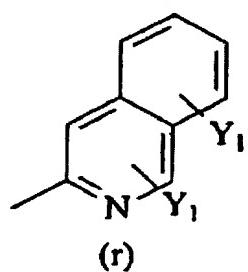
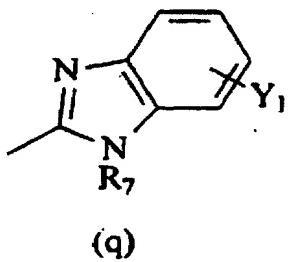
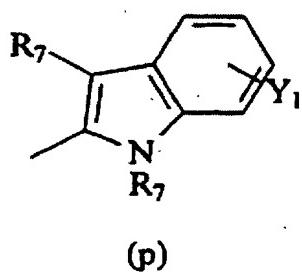
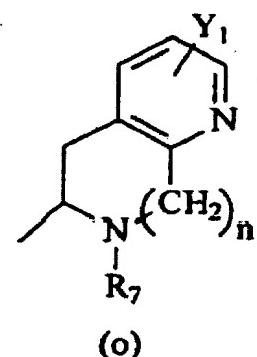
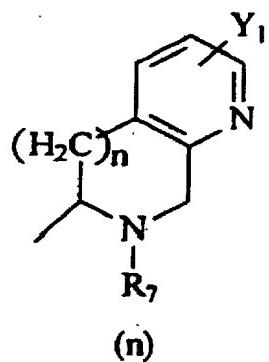
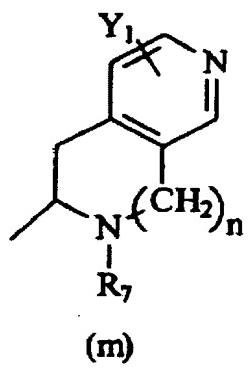
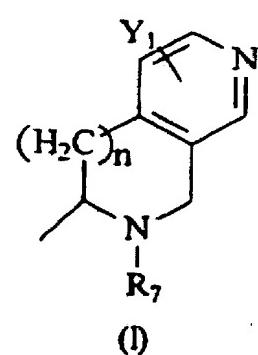
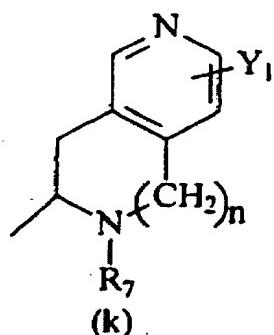
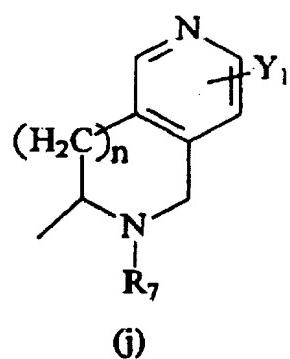
(g)

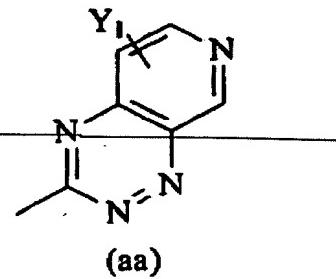
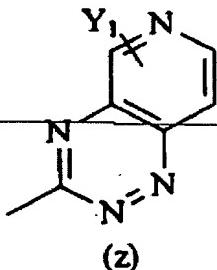
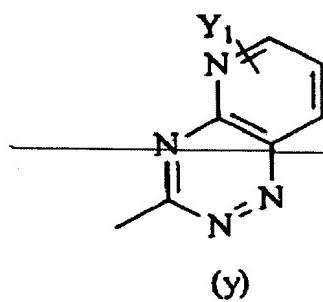
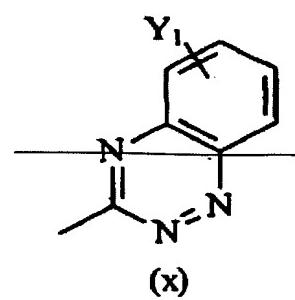
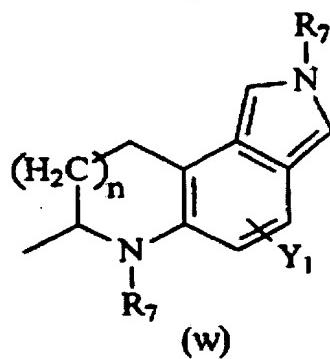
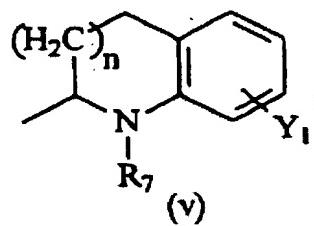
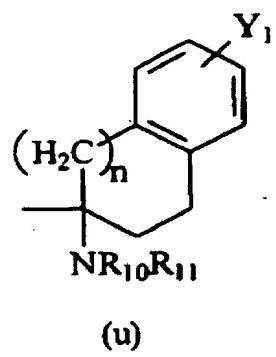
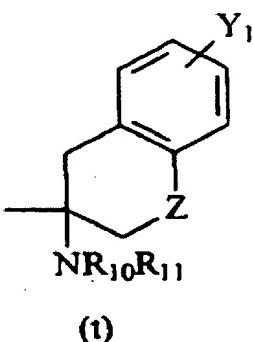
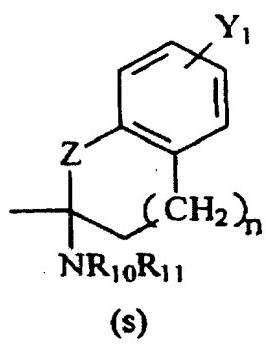


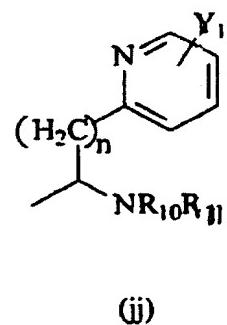
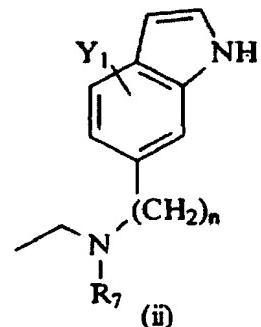
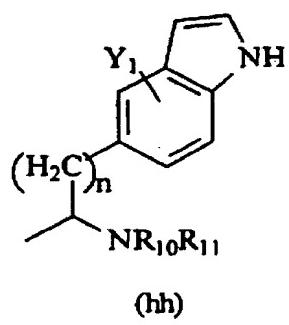
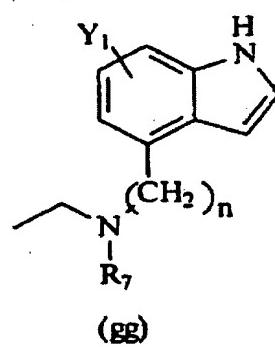
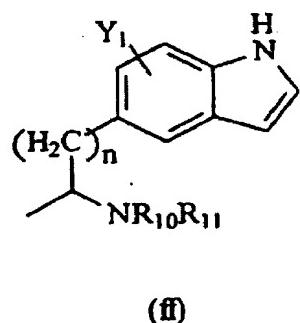
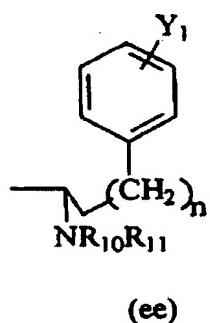
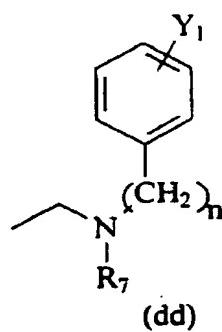
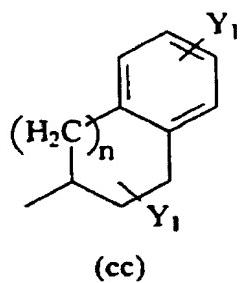
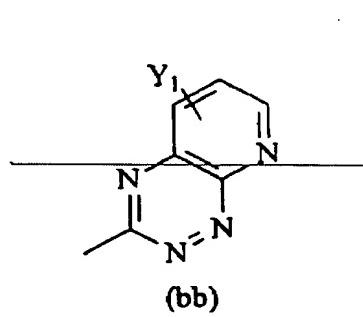
(h)

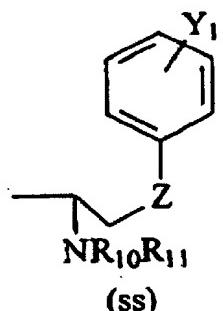
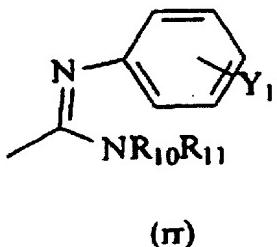
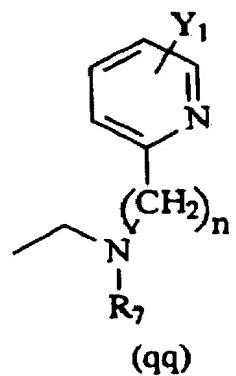
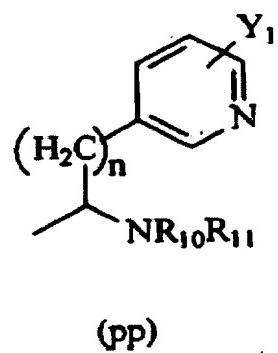
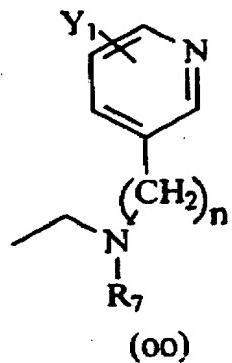
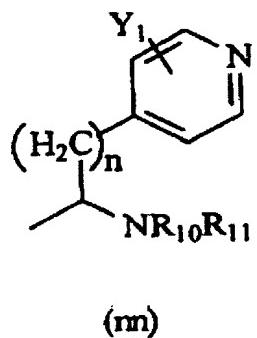
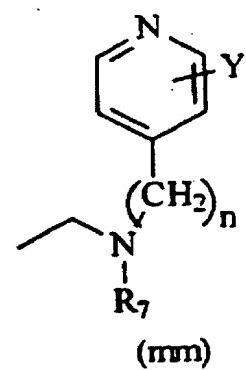
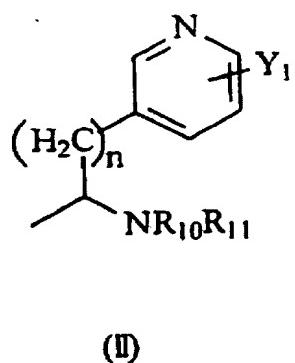
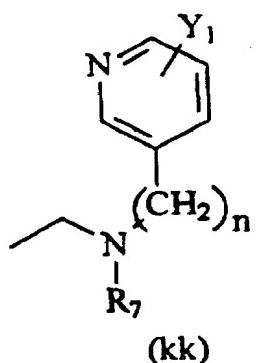


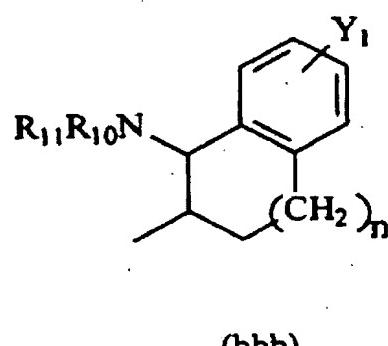
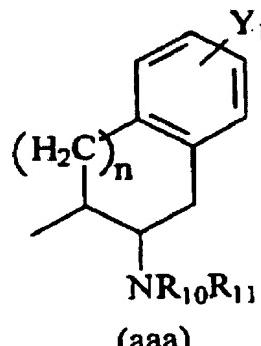
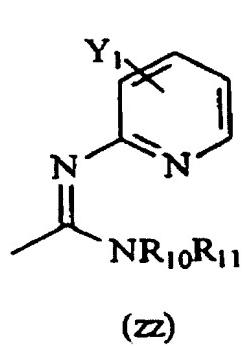
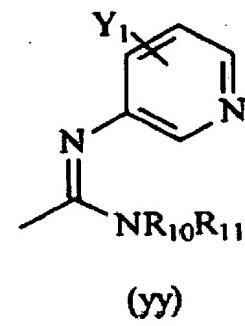
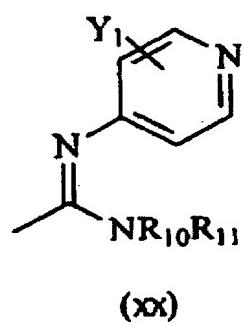
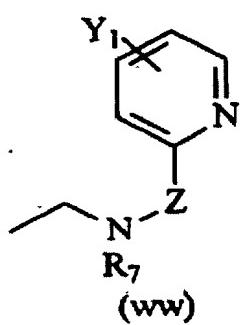
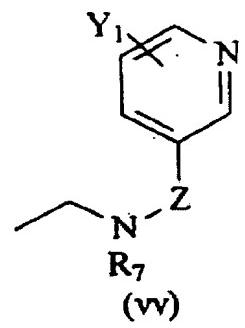
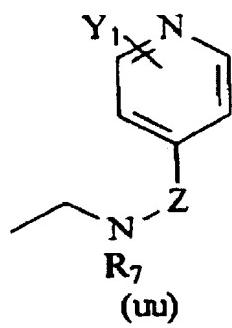
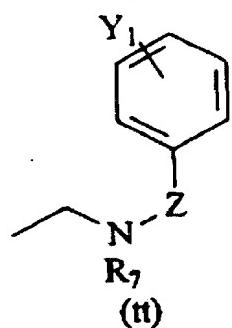
(i)











X₁ is hydrogen, C₁₋₈ alkyl, C₃₋₈alkenyl, or C₃₋₈alkynyl;

X₂ is hydrogen, C₁₋₈alkyl, C₃₋₈alkenyl, or C₃₋₈alkynyl;

or X₁ and X₂ together form =O, =S, =NH;

R₇ is H, C₁₋₈alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁,

NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, CH₂(CH₂)_nY₂, or C(=NH)NR₁₆R₁₇;

R₈ is H, C₁₋₈alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl;

R₉ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl;

R₁₀ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl ;

R₁₁ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl ;

R₁₂ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl ;

R₁₃ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl ;

R₁₄ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl ;

R₁₅ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl ;

R₁₆ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl ; and

R₁₇ is H, C₁₋₈ alkyl, CH₂ aryl substituted by one or more substituents H, OH, Br, Cl, F,

CN, CF₃, NO₂, N₃, C₁₋₆ alkyl, or CH₂(CH₂)_nY₂'; wherein Y₂' is H, CF₃, or C₁₋₆alkyl

or a pharmaceutically acceptable salt thereof.

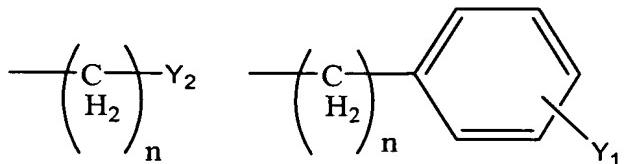
Claim 14. (Currently Amended) The pharmaceutical composition of claim 13, wherein said kappa opioid receptor antagonist is a compound of formula (I), wherein R₁, R₄, R₅, Y₁, Y₂, Z, n, X₁, X₂, and R₇-R₁₇ are as in Claim 13;

Y₃ is H;

R₂ and R₃ are each, independently, H, C₁₋₈ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, or CH₂aryl substituted by one or more substituents Y₁; and R₆ is a group having a formula selected from the group consisting of structures (a)-(w) and (cc).

Claim 15. (Previously Presented) The pharmaceutical composition of claim 13, wherein said kappa opioid receptor antagonist is a compound of formula (1), wherein Y₁, Y₂, R₄, R₅, Z, n, X₁, X₂ and R₈-R₁₅ are as in Claim 13;

R₁ is C₁₋₈ alkyl, or one of the following structures:



Y₃ is H;

R₂ and R₃ are each, independently, H or C₁₋₈ alkyl, wherein R₂ and R₃ cannot both be H at the same time;

R₆ is a formula selected from the structures (a)-(r) shown above; and

R₇ is H, C₁₋₈ alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, or CH₂(CH₂)_nY₂.

Claim 16. (Previously Presented) The pharmaceutical composition of claim 13, wherein said kappa opioid receptor antagonist is a compound of formula (I), wherein Y₁, Z, n, X₁, X₂ and R₈-R₁₅ are as noted- above in Claim 13;

R₁ is C₁₋₈ alkyl;

Y₂ is H, CF₃, CO₂R₉, C₁₋₆ alkyl, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, CH₂OH, CH₂OR₈, or COCH₂R₉;

Y₃ is H;

R₂ and R₃ are each, independently, H or methyl, wherein R₂ and R₃ cannot both be H at the same time;

R₄ is H, C₁₋₈ alkyl, CO₂C₁₋₈alkyl, or CH₂ aryl substituted by one or more substituents Y₁ and the stereocenter adjacent to R₄ is in an (S) configuration;

R₅ is H, C₁₋₈ alkyl, CH₂CO₂C₁₋₈ alkyl;

R₆ is a group having a formula selected from the group consisting of structures (a)-(c) and (h)-(o); and

R₇ is H, C₁₋₈alkyl, CH₂aryl substituted by one or more substituents Y₁, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₃, CONR₁₄R₁₅, or CH₂(CH₂)_nY₂.

Claim 17. (Previously Presented) The pharmaceutical composition of claim 13, wherein said kappa opioid receptor antagonist is a compound of formula (1), wherein Y₁, Z, n, X₁, X₂ and R₈-R₁₄ are as in Claim 13;

R₁ is methyl,

Y₂ is H, CF₃, CO₂R₉, C₁₋₆ alkyl, NR₁₀R₁₁, NHCOR₁₂, NHCO₂R₁₂, CONR₁₃R₁₄, CH20H, CH₂OR₈, or COCH₂R₉;

Y₃ is H;

R₂ and R₃ are each H or methyl, such that when R₂ is H, R₃ is methyl and vice versa;

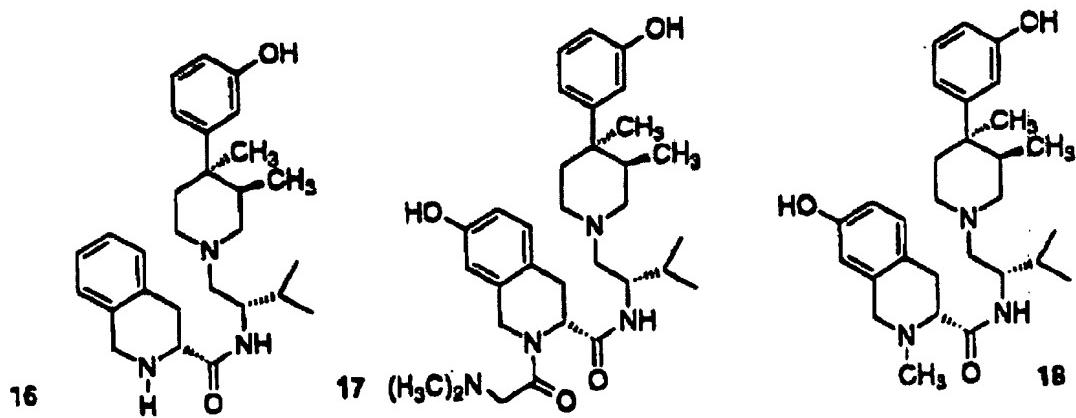
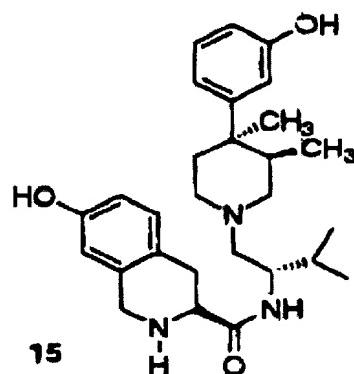
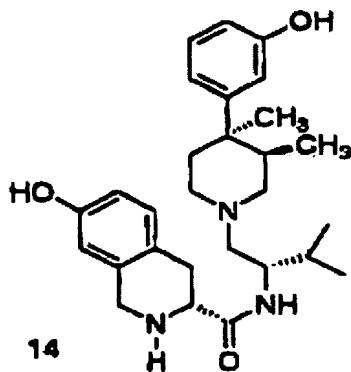
R₄ is C₁₋₈ alkyl, or CO₂C₁₋₈ alkyl, and the stereocenter adjacent to R₄ has a configuration of (S);

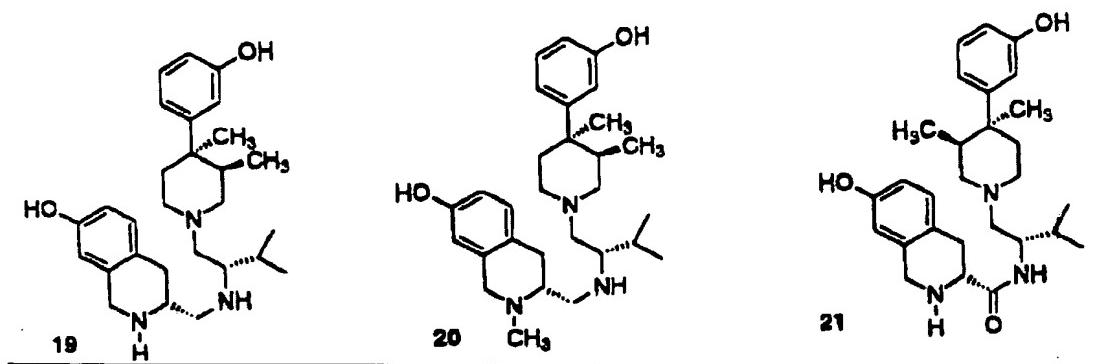
R₅ is H;

R₆ is a group having a formula selected from the group consisting of structures (a) and (b); and

R₇ is H, C₁₋₈ alkyl, CH₂aryl substituted by one or more substituents Y, or CH₂(CH₂)_nY₂.

Claim 18. (Currently Amended) The pharmaceutical composition of claim 13, wherein said kappa opioid receptor antagonist is a compound selected from formulae 14-21 of Fig. 1 as follows:



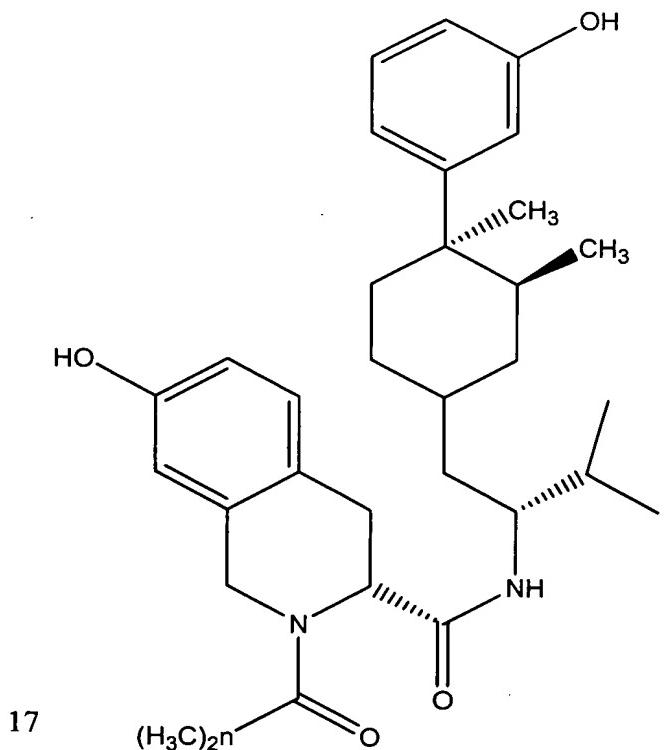


Claim 19. (Original) The pharmaceutical composition of claim 13, wherein said composition is an injectable composition.

Claim 20. (Original) The pharmaceutical composition of claim 13, wherein said composition is an orally administrable composition.

Claim 21. (Original) The pharmaceutical composition of claim 20, wherein said orally administrable composition is in a form selected from the group consisting of tablets, capsules, troches, powders, solutions, dispersions, emulsions and suspensions.

Claim 22. (Previously Presented) The kappa opioid receptor antagonist according to Claim 7, having the chemical formula:



Claim 23. (Cancelled)

Claim 24. (Cancelled)

Claim 25. (Cancelled)